WHAT IS CLAIMED:

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- 1. A low-temperature process for producing pigment-grade rutile titanium dioxide from an aqueous solution comprising the following sequential steps:
 - a. preparing an aqueous feed solution comprising a titanium compound;
 - b. adding an effective amount of a catalyzing salt to the solution;
 - c. optionally adding a chemical control agent to the solution;
- d. evaporating the solution to produce a dry amorphous intermediate that includes a mixture of titanium compounds; and
 - e. calcining the intermediate to form TiO₂ rutile pigment base.
- 10 2. The process of claim 1 further comprising washing the salt from the calcined TiO₂ rutile pigment base.
 - 3. The process of claim 2 further comprising milling and dispersing the TiO₂ rutile pigment.
- 4. The process of claim 1 wherein the titanium compound is selected from thegroup of titanium chloride, titanium oxychloride, and mixtures thereof.
 - 5. The process of claim 1 wherein the titanium compound is titanium oxychloride.
 - 6. The process of claim 1 wherein the catalyzing salt is a salt of an alkali metal.
 - 7. The process of claim 1 wherein the catalyzing salt is selected from the group consisting of chloride salts.
- 20 8. The process of claim 7 wherein the chloride salts are selected from the group consisting of NaCl, KCl, LiCl and mixtures thereof.
 - 9. The process of claim 7 wherein the chloride salts comprise a eutectic mixture of NaCl, KCl, and LiCl.

- 10. The process of claim 7 wherein the chloride salts comprise a eutectic mixture of LiCl and KCl.
- 11. The process of claim 7 wherein the chloride salts comprise a eutectic mixture of LiCl and NaCl.
- 5 12. The process of claim 1 wherein the catalyzing salt is present in the feed solution in an amount from about 3 weight % of the equivalent amount of TiO₂ present in the feed solution and the amount corresponding to the saturation point of the catalyzing salt in the feed solution.
- 13. The process of claim 1 wherein the catalyzing salt present in the feed solution is
 10 from about 10 weight % and about 50 weight % of the equivalent amount of TiO₂
 present in the feed solution.
 - 14. The process of claim 1 wherein the amount of the catalyzing salt is between about 15 weight % and about 30 weight % of the equivalent amount of TiO₂ present in the feed solution.
- 15. The process of claim 1 wherein the catalyzing salt does not significantly chemically react with titanium oxide through the process.
 - 16. The process of claim 1 wherein the catalyzing salt does not significantly change chemical composition.
 - 17. The process of claim 1 wherein the catalyzing salt is recycled.
- 20 18. The process of claim 1, wherein the TiO₂ rutile pigment base comprises an open network of rutile crystals.
 - 19. The process of claim 1 wherein the catalyzing salt has a melting point of less than 800°C.
- 20. The process of claim 1 wherein the calcining is conducted at a temperature less than 800°C.

- 21. The process of claim 1 wherein the calcining is conducted at a temperature less than 700°C.
- 22. The process of claim 1 wherein the calcining is conducted at a temperature less than 600°C.
- 5 23. The process of claim 1 wherein the calcining is conducted at a temperature less than 500°C.
 - 24. The process of claim 1 wherein the calcining is conducted at a temperature less than 400°C.
- 25. The process of claim 1 wherein the calcination time is between the time neededto melt the catalyzing salt and about 24 h.
 - 26. The process of claim 1 wherein the calcination time is less than about two hours.
 - 27. The process of claim 1 wherein the calcination time is less than about 30 minutes.
- 28. The process of claim 1 wherein the calcination time for is less than about one minute.
 - 29. The process of claim 1 wherein the chemical control agent is added before evaporating.
 - 30. The process of claim 27 wherein the chemical control agent is a water-soluble salt of tin.
- 20 31. The process of claim 27 wherein the chemical control agent is tin chloride.
 - 32. The process of claim 1 wherein the evaporating is conducted in a spray drier.
 - 33. The process of claim 1 wherein the evaporation temperature is between about 100°C and about 300°C.

- 34. The process of claim 1 wherein the amorphous intermediate comprises a homogeneous mixture of titanium, oxygen, chlorine, and hydrogen compounds, with a homogeneous distribution of salts through the titanium intermediate.
- 35. The process of claim 32 wherein the product after calcination comprises TiO₂ rutile crystallites bound in a structure of hollow spheres or parts of spheres.
 - 36. The process of claim 33 wherein the spheres have a diameter of about 0.1 to about 100 μ m.
 - 37. The process of claim 33 wherein the crystallites have a particle size between about 10 nm and 1000 nm.
- 10 38. The process of claim 33 wherein the crystallites have a particle size between about 50 nm and 500 nm.
 - 39. The process of claim 33 wherein the crystallites forming the hollow spheres have a particle size between about 100 nm and 300 nm.
- 40. The process of claim 1 wherein the washing is conducted with water to provide 15 an aqueous salt solution and the TiO₂ rutile pigment base.
 - 41. The process of claim 40 further comprising recycling the salts in the aqueous salt solution.
 - 42. The process of claim 41 further comprising milling the washed TiO₂ rutile pigment base product.
- 20 43. The process of claim 1 wherein a thermodynamically unstable brookite phase is formed as an intermediate during the early stages of calcination.